

TETRA-DELTA

100/105/100/100/10 mg,
intramamarna suspenzija, za
krave u laktaciji

Not
authorised

- Prednisolone
- Dihydrostreptomycin
- Benzylpenicillin procaine
- Neomycin
- Novobiocin

Product identification

Medicine name:

TETRA-DELTA 100/105/100/100/10 mg, intramamarna suspenzija, za krave u laktaciji

Active substance:

Prednisolone

Dihydrostreptomycin

Benzylpenicillin procaine

Neomycin

Novobiocin

Target species:

Cattle (lactating cow)

Route of administration:

Intramammary use

Product details

Active substance and strength:

Prednisolone

10.00 milligram(s) / 1.00 Syringe

Dihydrostreptomycin

100.00 milligram(s) / 1.00 Syringe

Benzylpenicillin procaine

100.00 milligram(s) / 1.00 Syringe

Neomycin

105.00 milligram(s) / 1.00 Syringe

Novobiocin

100.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary suspension

Withdrawal period by route of administration:**Intramammary use:**

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Cattle (lactating cow)

- Meat and offal. 7 day

- Milk. 108 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51RV01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Revoked

Authorised in:

Croatia

Package description:

Available only in [Croatian](#)

Available only in [Croatian](#)

Available only in [Croatian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Zoetis B.V.

Marketing authorisation date:

3/12/2020

Manufacturing sites for batch release:

Norbrook Laboratories Limited

Zoetis Belgium

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/20-01/751

Date of authorisation status change:

8/04/2022

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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